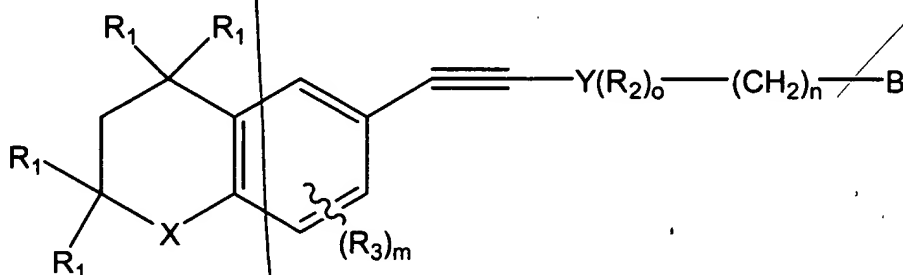


IN THE CLAIMS:

Amend Claims 1 and 14 as follows:

**Claim 1 (amended)**

A2  
A pharmaceutical composition for the treatment of a malignant disease or condition in a mammal, the composition comprising a pharmaceutically acceptable excipient and a therapeutically effective dose of a compound of the formula



Pub B1  
where X is S or O;

R<sub>1</sub> is, independently, H or lower alkyl of 1 to 6 carbons;

R<sub>2</sub> and R<sub>3</sub> are, independently, H, lower alkyl of 1 to 6 carbons, F, Cl, Br, I, alkoxy of 1 to 6 carbons, or fluoroalkoxy of 1 to 6 carbons;

m is an integer 0 to 3;

o is an integer 0 to 4;

n is 0-5;

Y is phenyl, naphthyl, or a heteroaryl group selected from a group consisting of pyridyl, thienyl, furyl, pyridazinyl, pyrimidinyl, pyrazinyl; oxazolyl, thiazolyl, or imidazolyl; and

B is COOH, a pharmaceutically acceptable salt thereof, CONR<sub>6</sub>R<sub>7</sub> or COOR<sub>8</sub> where R<sub>6</sub> and R<sub>7</sub>, independently, are hydrogen or an alkyl group of 1 to 6 carbons and R<sub>8</sub> is alkyl of 1 to 6 carbons,

said composition being adapted to be used in combination with another chemotherapeutic agent effective for the treatment of the malignant disease or

A2  
cont.

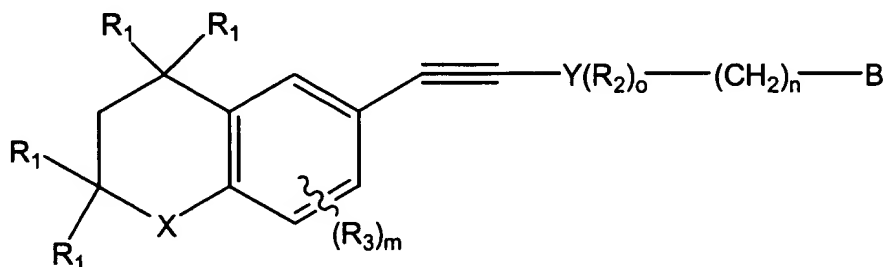
condition of the mammal where the composition in combination with the other chemotherapeutic agent shows synergistic effect.

#### Claim 14 (amended)

A3

A method of treating a malignant disease or condition in a mammal in need of such treatment, the method comprising the steps of:

administering to said mammal a pharmaceutical composition comprising a pharmaceutically acceptable excipient and a therapeutically effective dose of a compound of the formula



where **X** is S or O;

**R<sub>1</sub>** is, independently, H or lower alkyl of 1 to 6 carbons;

**R<sub>2</sub>** and **R<sub>3</sub>** are, independently, H, lower alkyl of 1 to 6 carbons, F, Cl, Br, I, alkoxy of 1 to 6 carbons, or fluoroalkoxy of 1 to 6 carbons;

**m** is an integer 0 to 3;

**o** is an integer 0 to 4;

**n** is 0-5;

**Y** is phenyl, naphthyl, or a heteroaryl group selected from a group consisting of pyridyl, thienyl, furyl, pyridazinyl, pyrimidinyl, pyrazinyl; oxazolyl, thiazolyl, or imidazolyl;

**B** is COOH, a pharmaceutically acceptable salt thereof, CONR<sub>6</sub>R<sub>7</sub> or COOR<sub>8</sub> where **R<sub>6</sub>** and **R<sub>7</sub>**, independently, are hydrogen or an alkyl group of 1 to 6 carbons and **R<sub>8</sub>** is alkyl of 1 to 6 carbons, and